Noromectin 10 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Ivermectin

Product identification

Medicine name:
Noromectin 10 mg/ml Solution for Injection for Cattle and Pigs

Active substance:
Ivermectin

Target species:
Cattle
Pig

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ivermectin
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 49 day
- **Pig**
  - Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

The product will be supplied in 1 L volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 500 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 250 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 100 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.
The product will be supplied in 50 ml volume, presented in high-density polyethylene vial with bromobutyl bung and aluminium cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
2/05/2001

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Laboratories (Ireland) Limited

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
51357

Date of authorisation status change:
31/10/2025

Reference member state:
Ireland

Procedure number:
IE/V/0104/001

Concerned member states:
France Germany Greece Iceland Italy Netherlands Portugal Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)
Published on: 29/12/2024
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